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§18-507.

- (a) The Department may, in consultation with the Statewide Steering Committee on Services for Adults with Sickle Cell Disease, provide services relating to sickle cell disease, including:
- (1) Educational programs on sickle cell disease for individuals affected by the disease, including:
 - (i) Individuals with sickle cell disease;
 - (ii) Families of individuals with sickle cell disease;
 - (iii) Caregivers of individuals with sickle cell disease;
 - (iv) Employees at primary and secondary schools; and
 - (v) Health care providers;
- (2) Social services support to individuals with sickle cell disease, including support from social workers and community health workers to provide information on services that may be available to the individual;
 - (3) Testing;
 - (4) Genetic counseling;
- (5) Assistance with any available reimbursement for medical expenses related to sickle cell disease;
- (6) Education and counseling services after the receipt of sickle cell trait test results from the State's Newborn Screening Program; and
- (7) Any other programs or services that are necessary to decrease the use of acute care services by individuals who have sickle cell disease.
- (b) The Department shall provide the services in subsection (a) of this section through community—based organizations to the extent practicable.

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